Address for all correspondence Research Ethics and Governance Office Royal Prince Alfred Hospital



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REFERENCE: X22-0396 & 2022/ETH02591

24 February 2023

This letter constitutes ethical approval only. You must NOT commence this research project at ANY site until you have submitted a Site Specific Assessment Form to the Research Governance Officer and received separate authorisation from the Chief Executive or delegate of that site.

Dear Professor Maguire,

Re: Protocol no. X22-0396 & 2022/ETH02591 - "An Online Skills-Based Educational Tool For Eating Disorders: SkillED Program Evaluation"

The Executive of the Ethics Review Committee, at its meeting of 24 February 2023 considered your correspondence of 14 February 2023. In accordance with the decision made by the Ethics Review Committee, at its meeting of 14 December 2022, ethical approval is granted.

I am pleased to advise that final ethical approval has been granted on the basis of the following:

• The research project meets the requirements of the National Statement on Ethical Conduct in Human Research.

This approval includes the following:

- HREA (Version 2, 14 February 2023)
- Protocol (Version 3, 20 February 2023)
- Participant Information Sheet Adult (Version 2, 14 February 2023)
- Participant Information Sheet Adolescent & Young Adult (Version 2, 14 February 2023)
- Participant Information Sheet Parent & Guardian (Version 2, 14 February 2023)
- Invitation and Consent Form- Adolescent/Young Adult & Parent/Guardian (Version 2, 14 February 2023)
- Invitation and Consent Form- Adult (Version 2, 14 February 2023)

- GP Clearance Form (Version 1, 14 February 2023)
- GP Discharge Letter (Version 1, 14 February 2023)
- Social Media Advertising (Version 3, 20 February 2023)
- Data Dictionary (Version 2, 14 February 2023)
- Disclaimer and Alert Messages (Version 3, 20 February 2023)
- Study Emails (Version 1, 17 November 2022)
- T0 Screening Questionnaire (Version 2, 14 February 2023)
- T1 Baseline Questionnaire (Version 2, 14 February 2023)
- T3 Completion of Core Modules (Version 2, 14 February 2023)
- T5 Completion of Optional Modules (Version 2, 14 February 2023)
- T6 3 Month Follow Up (Version 2, 14 February 2023)
- Weekly Monitoring Questionnaire (Version 1, 14 February 2023)
- SLHD Privacy Compliance Form (Version 3, October 2020)
- Research Data Management Plan (16 February 2023)
- T2 and T4 Before and after each module (undated)
- Semi Structured Interview (undated)
- Reasons For Withdrawal (undated)
- GCP Certificate Dr S Maguire (dated 21 October 2019)
- GCP Certificate S Barakat (dated 20 May 2022)
- GCP Certificate S Touyz (dated 27 January 2023)
- GCP Certificate D Rogers (dated 09 December 2022)
- GCP Certificate Marcellinus Kim (dated 07 February 2023)
- GCP Certificate M Pehlivan (dated 13 February 2023)
- GCP Certificate P Marks (dated 22 December 2022)
- GCP Certificate S Corry (dated 12 February 2023)
- GCP Certificate S H Ong (dated 09 January 2023)
- GCP Certificate J Miskovic-Wheatley (dated 13 February 2023)

- GCP Certificate K Spielman (dated 30 December 2022)
- Curriculum Vitae of Dr S Maguire
- Curriculum Vitae of Marcellinus Kim

You are asked to note the following:

The Committee noted that authorisation will be sought to conduct the study at the following sites:

- InsideOut Institute (eClinic platform)
- Sydney Local Health District
- It is a requirement of ethics approval that, before its commencement, this clinical trial is registered on a publicly accessible register, such as the Australian New Zealand Clinical Trials Registry or another appropriate international register. The Committee therefore sought details of the Register in which the study has been included and its registration number.
- This approval is valid for **five years**, and the Committee requires that you furnish it with annual reports on the study's progress beginning in **February 2024**. This will be through the submission of a milestone in REGIS.
- This human research ethics committee (HREC) has been accredited by the NSW Department of Health as a lead HREC under the model for single ethical and scientific review and is constituted and operates in accordance with the National Health and Medical Research Council's National Statement on Ethical Conduct in Human Research and the CPMP/ICH Note for Guidance on Good Clinical Practice.
- In accordance with the National Statement, Chapter 4.7, you are reminded that you must seek ethical approval from the HREC of the Aboriginal Health and Medical Research Council (AHMRC) if you intend to use Aboriginal or Torres Strait Islander status in any presentation or publication.
- **Partnering with Consumers:** As per Standard 2 of The National Clinical Trials Governance Framework, you are asked to provide an annual update with your annual progress report (milestone) on the ongoing involvement of consumers in the planning, design, delivery, measurement and evaluation of the trial.
- Good Clinical Practice (GCP): When adding additional sites, it is a condition
  of approval that the GCP Certificate of Completion be submitted for the
  principal investigator responsible for the new site.
- You must immediately report anything which might warrant review of ethical approval of the project in the specified format, including unforeseen events that might affect continued ethical acceptability of the project.
- You must notify the HREC of proposed changes to the research protocol or conduct of the research in the specified format.

 You must notify the HREC and other participating sites, giving reasons, if the project is discontinued at a site before the expected date of completion.

Should you have any queries about the Committee's consideration of your project, please contact me. The Committee's Terms of Reference, Standard Operating Procedures, membership and standard forms are available from the Sydney Local Health District website.

If you are not using REGIS, a copy of this letter must be forwarded to all site investigators for submission to the relevant Research Governance Officer.

The Ethics Review Committee wishes you every success in your research.

Regards,

Sanaa Thomas

P.Thomas

**Executive Officer** 

**Clinical Trials Sub-committee** 

For:

Rosemary Carney Executive Officer Ethics Review Committee (RPAH Zone)

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